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OM protein - protein search, using sw model

Run on: December 30, 2002, 16:14:27 ; Search time 35 Seconds

(without alignments)  
247.465 Million cell updates/sec

Title: US-09-664-326-23

Perfect score: 368  
Sequence: 1 LVTDCESGQNICLCEGSN.....KPFOSHNDGFEEPEEYIQ 65

Scoring table:

BL0SUM62  
Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%  
Listing first 45 summaries

Database :

1: /SIDSM2/gcgdata/geneseq/geneseq-emb1/AA1980.DAT:\*  
2: /SIDSM2/gcgdata/geneseq/geneseq-emb1/AA1981.DAT:\*  
3: /SIDSM2/gcgdata/geneseq/geneseq-emb1/AA1982.DAT:\*  
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19: /SIDSM2/gcgdata/geneseq/geneseq-emb1/AA1998.DAT:\*  
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22: /SIDSM2/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:\*  
23: /SIDSM2/gcgdata/geneseq/geneseq-emb1/AA2002.DAT:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	368	100.0	65	10	AAp90359	Hirudin derivative
2	368	100.0	65	16	AAp78291	Desulphatohirudin
3	368	100.0	65	16	AAp79813	Hirudin derivative
4	368	100.0	65	17	AAW13897	Hirudin variant (L
5	368	100.0	65	17	AAW03735	Recombinant hirudi
6	368	100.0	65	18	AAW11527	S. marcescens hiru
7	368	100.0	65	22	AAp70828	Hir(1-65)/AspPro/T
8	368	100.0	127	23	AAp80818	Hirudin variant (d
9	365	99.2	65	17	AAW13896	Anticoagulant pept
10	364	98.9	64	6	AAp50082	

11	363	98.6	64	15	AAp59773	Desulphatohirudin.
12	360	97.8	65	6	AAp50329	Hirudin protein.
13	360	97.8	65	6	AAp50335	Hirudin variant.
14	360	97.8	65	6	AAp50188	Desulphatohirudine
15	360	97.8	65	8	AAp70225	Sequence of desulp
16	360	97.8	65	12	AAp12887	Synthetic hirudin
17	360	97.8	65	13	AAp24308	Hirudin HV-1. Syn
18	360	97.8	65	14	AAp30486	Hirudin HV1. Hiru
19	360	97.8	65	15	AAp54636	Anticoagulant hiru
20	360	97.8	65	15	AAp59767	Desulphatohirudin
21	360	97.8	65	16	AAp62214	HV1 derivative of
22	360	97.8	65	16	AAp69029	Hirudin variant 1
23	360	97.8	65	17	AAp9354	Hirudin variant HV
24	360	97.8	65	17	AAW13889	Hirudin variant (I
25	360	97.8	65	18	AAW21762	Hirudin, fused to
26	360	97.8	65	19	AAp82265	Hirudin variant RH
27	360	97.8	65	23	AAp77700	Hirudin amino acid
28	360	97.8	66	7	AAp60395	Desulphatohirudin.
29	360	97.8	66	12	AAp10969	HV-1. Synthetic.
30	360	97.8	66	13	AAp31209	Desulphatohirudin
31	360	97.8	66	13	AAp31210	Desulphatohirudin
32	360	97.8	66	17	AAW13892	Hirudin variant (P
33	360	97.8	69	13	AAp24309	Hirudin (HV-1) RGD
34	360	97.8	69	13	AAp24312	Hirudin (HV-1) RGD
35	360	97.8	70	13	AAp24315	Hirudin (HV-1) GRE
36	360	97.8	71	13	AAp24327	Hirudin (HV-1) GRG
37	360	97.8	72	13	AAp24321	Hirudin (HV-1) CRG
38	360	97.8	72	13	AAp24324	Hirudin (HV-1) CRG
39	360	97.8	77	13	AAp24318	Hirudin (HV-1) HHL
40	360	97.8	82	15	AAp54088	PHOS leader and hl
41	360	97.8	82	15	AAp51073	yeast PHOS signal
42	360	97.8	92	12	AAp14151	MSP signal peptide
43	360	97.8	93	15	AAp47489	HV-1 encoded by su
44	360	97.8	93	15	AAp47490	Desulphatohirudin
45	360	97.8	134	12	AAp12888	Factor Xa-cleavabl

ALIGNMENTS

RESULT 1	
AAp90359	
ID	AAp90359 standard; protein: 65 AA.
XX	
AC	AAp90359;
XX	
DT	01-NOV-1989 (first entry)
XX	
DE	Hirudin derivative.
XX	
KW	Hirudin deriv; thrombin inhibitor.
PN	
XX	EP324712-A.
XX	
PD	19-JUL-1989.
XX	
PF	13-JAN-1988; 88EP-080540.
XX	
PR	13-JAN-1988; 88DE-380540.
XX	
PA	(FARH ) HOECHST AG.
XX	
PI	Crause P, Habermann P, Tripiet D;
DR	WPL; 1989-208655/29.
XX	
PT	New hirudin deriv. with N-terminal leucine - is expressed in high
XX	yields in yeasts and is secreted in form with correct folding.
PS	Claim 1; page 8; 11pp; German.
XX	
CC	The hirudin deriv. has thrombin-inhibiting activity. Unlike
CC	analogues with N-terminal Thr-Tyr or Ile-Tyr units, it is

CC expressed in high yields in yeasts and is secreted in a  
 CC form with correct folding.

XX Sequence 65 AA;

Query Match 100.0%; Score 368; DB 10; Length 65;  
 Best Local Similarity 100.0%; Pred. No. 1.8e-28;  
 Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 LVTYDCTESGQNLCEGSSNVCQGKNCITLSDGKKNOCVTGEGTPKPSHNDGDFEELP 60  
 DB 1 LVTYDCTESGQNLCEGSSNVCQGKNCITLSDGKKNOCVTGEGTPKPSHNDGDFEELP 60  
 OY 61 EYLIQ 65  
 DB 61 EYLIQ 65

#### RESULT 2

AAR78291  
 ID AAR78291 standard; protein; 65 AA.

AC AAR78291;

XX 06-MAR-1996 (first entry)

DE Desulphatohirudin HVL.

KW Desulphatohirudin; leech; Hirudo medicinalis; anticoagulant; sugar;  
 stability; therapy.

OS Hirudo medicinalis.

XX W09520399-A1.

PD 03-AUG-1995.

PF 25-JAN-1995; 95MO-1B00053.

XX 26-JAN-1994; 94GB-0001447.

PA (CIBA ) CIBA GEIGY AG.

PI Arvinte T;

DR WPI; 1995-275296/36.

XX New freeze dried hirudin compositions - contg. potassium phosphate  
 PT and a sugar to provide long term storage stability at ambient temps.

XX Disclosure; Page 3; 22pp; English.

CC The amino acid sequence of the desulphatohirudin composition HVL.  
 CC The hirudin cpds. AAR78290-4 can be isolated from leeches (Hirudo  
 CC medicinalis). The cpds. have anticoagulant properties and are  
 CC useful in compositions contg. the hirudin, potassium phosphate and  
 CC a sugar pref. mannitol, trehalose, sucrose, etc. The potassium  
 CC phosphate has been found to increase the stability of the hirudin  
 CC cpd. esp. at ambient temp. The comps. contg. the hirudin can be  
 CC used for anticoagulant therapy.

XX Sequence 65 AA;

Query Match 100.0%; Score 368; DB 16; Length 65;  
 Best Local Similarity 100.0%; Pred. No. 1.8e-28;  
 Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 LVTYDCTESGQNLCEGSSNVCQGKNCITLSDGKKNOCVTGEGTPKPSHNDGDFEELP 60  
 DB 1 LVTYDCTESGQNLCEGSSNVCQGKNCITLSDGKKNOCVTGEGTPKPSHNDGDFEELP 60  
 OY 61 EYLIQ 65  
 DB 61 EYLIQ 65

DB 61 EYLIQ 65

#### RESULT 3

AAR79813  
 ID AAR79813 standard; protein; 65 AA.

XX AAR79813;

XX 28-MAR-1996 (first entry)

DE Hirudin derivative.

KW Hirudin; derivative; anticoagulant; polyethylene glycol.

XX Synthetic.

PN EP667355-A1.

PD 16-AUG-1995.

PF 06-FEB-1995; 95EP-0101554.

XX 10-FEB-1994; 94DE-4404168.

PA (FARH ) HOECHST AG.

PI Hropot M, Ludwig J, Obermeier R, Tripler D;

DR WPI; 1995-276615/37.

XX New hirudin deriv. with amine deriv. attached to position 36 or 63  
 PT - useful as anticoagulants, partic. for transdermal delivery by  
 PT iontophoresis.

XX Disclosure; Page 8; 14pp; German.

CC Hirudin derivatives of formula A0-A1-A2-(Hirudin 3-36)-(Y)-(Hirudin  
 CC 37-65) have anticoagulant activity, especially those derivatised  
 CC with polyethylene glycol. In the formula A0, A1 and A2 are amino  
 CC acid residues and A0 can also be H, Y is a residue of amines NH2-R-X  
 CC or A-RI-X, where A is 1-10 amino acids, R is a 1-10C alkyl (opt.  
 CC substituted), RI is either H, a covalent bond, 1-10 sugar residues  
 CC or (O-(CH2)m)n where m is 2-5 and n is 1-100. X is H, OR2, NHR2, C  
 CC OR2 or an amino acid. R2 is H or as R. The - sign denotes that the  
 CC two hirudin fragments are connected by disulphide bridges.

XX Sequence 65 AA;

Query Match 100.0%; Score 368; DB 16; Length 65;  
 Best Local Similarity 100.0%; Pred. No. 1.8e-28;  
 Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 LVTYDCTESGQNLCEGSSNVCQGKNCITLSDGKKNOCVTGEGTPKPSHNDGDFEELP 60  
 DB 1 LVTYDCTESGQNLCEGSSNVCQGKNCITLSDGKKNOCVTGEGTPKPSHNDGDFEELP 60  
 OY 61 EYLIQ 65  
 DB 61 EYLIQ 65

#### RESULT 4

AAW13897  
 ID AAW13897 standard; Protein; 65 AA.

XX AAW13897;

XX 14-MAY-1997 (first entry)

DE Hirudin variant (Leu 1, Thr 2)-desulphato hirudin HVL.

KW Hirudin; variant; thrombin inhibitor; human; acetylsalicylic acid; ASA;

thrombolytic agent; cardiovascular event; stroke; cardiovascular death;  
 coronary re-vascularisation; therapy; acute myocardial infarction; AMI;  
 hirudo medicinalis.  
 Synthetic.  
 Key Location/Qualifiers  
 Misc-difference 1 /Label= V1L  
 Misc-difference 2 /Label= V2T  
 Modified-site 63 /note= "modified with phenolic hydroxy group"  
 PF 12-MAR-1996; 96EP-0103821.  
 PD 18-SEP-1996.  
 PR 12-MAY-1995; 950S-0440556.  
 PR 15-MAY-1995; 950S-0405269.  
 PA (BEHM ) BEHRINGER AG.  
 PA (BGM ) BRIGHAM & WOMENS HOSPITAL.  
 PI Heinrichs H, Hennekens CH;  
 DR WPI; 1996-414245/42.  
 XX Composition comprising acetyl:salicylic acid and hirudin - is esp.  
 PT useful for preventing the recurrence of acute myocardial  
 PT infarction(s)  
 Claim 6; : 11pp; English.  
 PS  
 XX  
 CC AAM13889-W13898 represent mutations of the hirudin variants represented  
 CC by AAR9354-R9356. Hirudin is a direct thrombin inhibitor, which has a  
 CC very high affinity for human (as well as other mammalian species)  
 CC thrombin. One molecule binds to a thrombin molecule, forming a tight  
 CC noncovalent complex and thereby irreversibly inactivates thrombin. These  
 CC sequences can be used in a composition of the invention, which also  
 CC contains acetyl:salicylic acid (ASA). The composition may be administered  
 CC to patients not undergoing treatment with a thrombolytic agent, to  
 CC inhibit and/or prevent myocardial or cardiovascular events (including  
 CC myocardial infarctions, strokes, coronary re-vascularisation or  
 CC cardiovascular death) in the patient. The compositions of the invention  
 CC are especially useful for preventing the recurrence of acute myocardial  
 CC infarctions (AMI). The components ASA and hirudin act synergistically.  
 CC The combined use of ASA and hirudin in AMI patients where thrombolytic  
 CC treatment is not advisable is expected to result in a higher incidence of  
 CC open coronary vessels.  
 CC  
 CC  
 SQ Sequence 65 AA;  
 Query Match 100.0%; Score 368; DB 17; Length 65;  
 Best Local Similarity 100.0%; Pred. No. 1.8e-28;  
 Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 1 LTYTDCESGONLCLCGSNVCGGKNCILGSDGKNCVYGEETPKQSHNDGFEETP 60  
 Db 1 LTYTDCESGONLCLCGSNVCGGKNCILGSDGKNCVYGEETPKQSHNDGFEETP 60  
 Oy 61 EYVLIQ 65  
 Db 61 EYVLIQ 65  
 RESULT 5  
 AAM03735  
 ID AAM03735 standard; protein; 65 AA.  
 XX  
 AC AAM03735;

17-OCT-1996 (first entry)  
 Recombinant hirudin analogue for admin. by intravenous drip injection.  
 DE  
 XX  
 KW Hirudin; anti-coagulant; disseminated intravascular coagulation; DIC;  
 KW thrombin inhibitor; low dosage; reduced side-effects; bleeding.  
 XX  
 OS Synthetic.  
 PN JP08143470-A.  
 PD 04-JUN-1996.  
 PF 18-NOV-1994; 94JP-0284910.  
 PR 18-NOV-1994; 94JP-0284910.  
 PA (FARH ) HOECHST JAPAN KK.  
 DR WPI; 1996-318859/32.  
 XX  
 PT Admin. of specific, lower dosage of hirudin or analogue by  
 PT intravenous drip injection - reduces side-effects in treatment of  
 PT disseminated intravascular coagulation  
 PS Claim 3; Page 2; 5pp; Japanese.  
 XX  
 CC The present sequence is that of the preferred hirudin analogue to be  
 CC administered in a novel intravenous drip injection for treatment of  
 CC disseminated intravascular coagulation. The hirudin molecule is  
 CC formulated at a concentration of 0.005-0.038 mg/ml (50-380 ATU/ml);  
 CC admin. of a reduced dosage of hirudin suppresses unwanted bleeding.  
 CC  
 CC  
 SQ Sequence 65 AA;  
 Query Match 100.0%; Score 368; DB 17; Length 65;  
 Best Local Similarity 100.0%; Pred. No. 1.8e-28;  
 Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 1 LTYTDCESGONLCLCGSNVCGGKNCILGSDGKNCVYGEETPKQSHNDGFEETP 60  
 Db 1 LTYTDCESGONLCLCGSNVCGGKNCILGSDGKNCVYGEETPKQSHNDGFEETP 60  
 Oy 61 EYVLIQ 65  
 Db 61 EYVLIQ 65  
 RESULT 6  
 AAM11527  
 ID AAM11527 standard; protein; 65 AA.  
 XX  
 AC AAM11527;  
 DT 11-SEP-1997 (first entry)  
 DE Recombinant hirudin derivative.  
 XX  
 KW hirudin; recombinant; derivative; treatment; prevention; brain tissue;  
 KW cellular infiltration; polynuclear leukocyte; monocyte; macrophage;  
 KW inhibit; vimentin positive astrocyte; anti-inflammatory.  
 XX  
 OS Synthetic.  
 PN JP08310967-A.  
 PD 26-NOV-1996.  
 PF 17-MAY-1995; 95JP-0118388.  
 PR 17-MAY-1995; 95JP-0118388.

PA (FARH ) HOECHST JAPAN LTD.  
XX  
DR WPI: 1997-061735/06.  
XX  
PT Agent for treatment and prevention of brain tissue damage -  
PT comprises hirudin or deriv. as active ingredient to inhibit damage  
PT caused by inflammatory cell infiltration  
XX  
PS Claim 3; Page 2; 5pp; Japanese.  
XX  
CC This sequence is a preferred recombinant hirudin derivative for use as  
CC an agent for treatment and prevention of brain tissue damage.  
CC particularly secondary damage caused by cellular infiltration of  
CC polynuclear leukocytes or the monocyte/macrophage system. The agent is  
CC effective against damage caused by inflammatory cells and inhibits the  
CC expression of vimentin positive astrocytes with high anti-inflammatory  
CC effect. Hirudin or its derivs. are used to prepare conventional  
CC pharmaceutical preps. for admin. by drip infusion or local injection  
CC at a dosage of about 0.001-5 mg/day for a male adult patient.  
XX  
SQ Sequence 65 AA;  
Query Match 100.0%; Score 368; DB 18; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.8e-28;  
Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 LRYTDCESGQNLCLCEGSNVCQGQNKCLIGSDGERKNCVTGEGTPKPSHNDGDFEIRP 60  
DB 1 LRYTDCESGQNLCLCEGSNVCQGQNKCLIGSDGERKNCVTGEGTPKPSHNDGDFEIRP 60  
QY 61 EYTLQ 65  
DB 61 EYTLQ 65  
RESULT 7  
AAB70828 ID AAB70828 standard; Protein: 65 AA.  
XX  
AC AAB70828;  
XX  
DT 18-JUN-2001 (first entry)  
XX  
DE S. marcescens hirudin protein fragment.  
XX  
KW Hirudin; Outer membrane protein; OPRF; lambd; fumarate reductase;  
KW Leu-Hirudin; Leu1-Thr2-63-desulfato-hirudin; antithrombotic.  
XX  
OS Serratia marcescens.  
XX  
PN DE1944870-A1.  
XX  
PD 29-MAR-2001.  
XX  
PF 18-SEP-1999; 99DE-1044870.  
XX  
PR 18-SEP-1999; 99DE-1044870.  
XX  
PA (AVET ) AVENTIS PHARMA DEUT GMBH.  
XX  
PI Habermann P, Bender R;  
XX  
DR WPI: 2001-246103/26.  
DR N-PSDB; AAF61507.  
XX  
XX Hirudin precursor containing heterologous signal peptide, useful for  
PT recombinant production of antithrombotic Leu-hirudin, is efficiently  
PT secreted and processed -  
XX  
PS Disclosure; Page 9; 12pp; German.  
XX  
CC This invention describes a novel hirudin precursor (I), comprising the  
CC signal sequence from the outer membrane protein of Serratia marcescens.

CC the OPRF protein of Pseudomonas fluorescens, the lambd protein of  
CC Escherichia coli, or the fumarate reductase of Shewanella putrefaciens,  
CC with the Leu-hirudin (LH) ((Leu1-Thr2)-63-desulfato-hirudin) sequence  
CC linked to the C-terminus of the signal sequence. (I) is an intermediate  
CC in recombinant production of LH, a known antithrombotic. The specified  
CC signal sequence may also be used for secretory expression of other  
CC proteins. (I) is processed directly to LH and this, in native form,  
CC secreted from E. coli in high yield. This results, both during  
CC fermentation and subsequent purification, in a higher concentration of  
CC hirudin, reducing costs of production. The specified signal sequences  
CC provide more efficient secretion than known sequences. This sequence  
CC represents a fragment of the S. marcescens hirudin protein.  
XX  
SQ Sequence 65 AA;  
Query Match 100.0%; Score 368; DB 22; Length 65;  
Best Local Similarity 100.0%; Pred. No. 1.8e-28;  
Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 LRYTDCESGQNLCLCEGSNVCQGQNKCLIGSDGERKNCVTGEGTPKPSHNDGDFEIRP 60  
DB 1 LRYTDCESGQNLCLCEGSNVCQGQNKCLIGSDGERKNCVTGEGTPKPSHNDGDFEIRP 60  
QY 61 EYTLQ 65  
DB 61 EYTLQ 65  
RESULT 8  
ABB08618 ID ABB08618 standard; Protein: 127 AA.  
XX  
AC ABB08618;  
XX  
DT 10-APR-2002 (first entry)  
XX  
DE Hir(1-65)/AspPro/TAP(1-60) fusion protein.  
XX  
KW Fusion protein; hirudin; Hir; tick anticoagulant protein; TAP;  
KW anticoagulant; blood; thrombin; factor Xa.  
XX  
OS Chimeric - Hirudo medicinalis.  
OS Chimeric - Ornithodoros moubata.  
OS Synthetic.  
XX  
FH Key Location/Qualifiers  
FT Region 1..65  
FT Region /label= hirudin  
FT Region 66..67  
FT /label= Asp/Pro-linker  
FT Region 68..127  
FT /label= TAP  
XX  
PN WO200204486-A2.  
XX  
PD 17-JAN-2002.  
XX  
PF 27-JUN-2001; 2001WO-EP07333.  
XX  
PR 07-JUL-2000; 2000DE-1033195.  
XX  
PA (AVET ) AVENTIS PHARMA DEUT GMBH.  
XX  
PI Habermann P;  
XX  
DR WPI: 2002-154918/20.  
DR  
XX New fusion protein of hirudin and tick anticoagulant protein, useful as  
PT an anticoagulant -  
XX  
PS Example 4; Page -; 36pp; German.  
XX  
CC The invention relates to a bifunctional fusion protein comprising

CC hirudin or its variant and tick anticoagulant protein (TAP) or its  
 CC variant. The fusion proteins have anticoagulant activity and are used to  
 CC inhibit coagulation of blood by acting to inhibit thrombin and inhibit  
 CC factor Xa. The present sequence is that of the Hir(1-65)/AspPro/TAP(1-60)  
 CC fusion protein useful in the illustration of the invention.  
 CC Note: The present sequence is not given in the specification but is  
 CC derived from SEQ ID NO 15 (ABB08603) and SEQ ID NO 17 (ABB08606).  
 XX  
 SO Sequence 127 AA;  
 Query Match 100.0%; Score 368; DB 23; Length 127;  
 Best Local Similarity 100.0%; Pred. No. 3,4e-28;  
 Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 LVTYDCTESGONLCICBSNVCGGKNCILGSDGKNCQVTEGTPKQSHNDGDFEIRP 60  
 Db 1 LVTYDCTESGONLCICBSNVCGGKNCILGSDGKNCQVTEGTPKQSHNDGDFEIRP 60  
 QY 61 EYVLIQ 65  
 Db 61 EYVLIQ 65  
 RESULT 9  
 AAM13896  
 ID AAM13896 standard; Protein; 65 AA.  
 XX  
 AC AAM13896;  
 XX  
 DT 14-MAY-1997 (first entry)  
 DE Hirudin variant (des-Val 1, Thr 2)-desulphato hirudin HVL.  
 XX  
 KW Hirudin; variant; thrombin inhibitor; human; acetylsalicylic acid; ASA;  
 KW thrombolytic agent; cardiovascular event; stroke; cardiovascular death;  
 KW coronary re-vascularisation; therapy; acute myocardial infarction; AMI;  
 KW hirudo medicinalis.  
 XX  
 OS Synthetic.  
 XX  
 FH Key Location/Qualifiers  
 FT Misc-difference 1 /note- "D-form residue"  
 FT Misc-difference 2 /label= V2T  
 FT Modified-site 63 /note- "modified with phenolic hydroxy group"  
 FT  
 XX  
 PN EP732102-A2.  
 XX  
 PD 18-SEP-1996.  
 XX  
 PF 12-MAR-1996; 96EP-0103821.  
 XX  
 PR 12-MAY-1995; 95US-0440556.  
 PR 15-MAR-1995; 95US-0405269.  
 XX  
 PA (BEHN ) BEHRINGWERKE AG.  
 PA (BGHM ) BRIGHAM & WOMENS HOSPITAL.  
 XX  
 PI Heinrichs H, Hennekens CH;  
 XX  
 DR WPI; 1996-414245/42.  
 XX  
 PT Composition comprising acetyl-salicylic acid and hirudin - is esp.  
 PT useful for preventing the recurrence of acute myocardial  
 PT infarction(s)  
 XX  
 PS Claim 6; ; 11pp; English.  
 XX  
 CC AAM13889-M13898 represent mutations of the hirudin variants represented  
 CC by AAB99354-R99356. Hirudin is a direct thrombin inhibitor, which has a  
 CC very high affinity for human (as well as other mammalian species)

CC thrombin. One molecule binds to a thrombin molecule, forming a tight  
 CC noncovalent complex and thereby irreversibly inactivates thrombin. These  
 CC sequences can be used in a composition of the invention, which also  
 CC contains acetylsalicylic acid (ASA). The composition may be administered  
 CC to patients not undergoing treatment with a thrombolytic agent, to  
 CC inhibit and/or prevent myocardial or cardiovascular events (including  
 CC myocardial infarctions, strokes, coronary re-vascularisation or  
 CC cardiovascular death) in the patient. The compositions of the invention  
 CC are especially useful for preventing the recurrence of acute myocardial  
 CC infarctions (AMI). The components ASA and hirudin act synergistically.  
 CC The combined use of ASA and hirudin in AMI patients where thrombolytic  
 CC treatment is not advisable is expected to result in a higher incidence of  
 CC open coronary vessels.  
 XX  
 SO Sequence 65 AA;  
 Query Match 99.2%; Score 365; DB 17; Length 65;  
 Best Local Similarity 98.5%; Pred. No. 3,4e-28;  
 Matches 64; Conservative 1; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 LVTYDCTESGONLCICBSNVCGGKNCILGSDGKNCQVTEGTPKQSHNDGDFEIRP 60  
 Db 1 LVTYDCTESGONLCICBSNVCGGKNCILGSDGKNCQVTEGTPKQSHNDGDFEIRP 60  
 QY 61 EYVLIQ 65  
 Db 61 EYVLIQ 65  
 RESULT 10  
 AAP50082  
 ID AAP50082 standard; Protein; 64 AA.  
 XX  
 AC AAP50082;  
 XX  
 DT 22-OCT-1991 (first entry)  
 DE Anticoagulant peptide.  
 XX  
 DE Anticoagulant peptide.  
 XX  
 KW Anticoagulant; diagnosis;  
 KW hirudo medicinalis.  
 XX  
 OS Hirudo medicinalis.  
 XX  
 PN EP158986-A.  
 PD 23-OCT-1985.  
 XX  
 PF 12-APR-1985; 85EP-0104445.  
 XX  
 PR 18-APR-1984; 84DE-3414593.  
 PR 19-OCT-1984; 84DE-3438296.  
 XX  
 PA (FARH ) HOECHST AG.  
 XX  
 PI Tripler D;  
 XX  
 DR WPI; 1985-264974/43.  
 XX  
 PT New polypeptide cpds. with anticoagulant activity - extracted from  
 PT leeches and synthetic analogues.  
 XX  
 PS Disclosure; page 2; 24pp; german.  
 XX  
 CC The peptide and its cleavage prods. are useful as anticoagulants. They  
 CC are specific stoichiometric inhibitors of thrombin, so can be used  
 CC therapeutically or as reagents for diagnosis. The C-terminal Tyr residue  
 CC has a phenolic H or phenol ester gp., Pref. H, SO3H or PO3H2.  
 XX  
 SO Sequence 64 AA;  
 Query Match 98.9%; Score 364; DB 6; Length 64;  
 Best Local Similarity 100.0%; Pred. No. 4,2e-28;  
 Matches 64; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 TTTDCTESGONCLCEGSNVCGGKNCILGSDGKNCQVTEGTPKPOSHNDGFEELP 61  
 DB 1 TTTDCTESGONCLCEGSNVCGGKNCILGSDGKNCQVTEGTPKPOSHNDGFEELP 60  
 OY 62 EYLIQ 65  
 DB 61 EYLIQ 64

## RESULT 11

AAR59773  
 ID AAR59773 standard; peptide; 64 AA.

AC AAR59773;

DT 17-FEB-1995 (first entry)

DE Desulphatohirudin.

XX Desulphatohirudin; variant; sulphate monoester group; hirudin;  
 KW depot formulation; deep vein thrombosis; water; calcium; magnesium;  
 KM zinc; ions; water-insoluble salt; stability; bleeding.

XX Hirudo medicinalis.

PN NZ250895-A.

PD 27-JUN-1994.

PF 16-FEB-1994; 94NZ-0250895.

PR 18-FEB-1993; 93GB-0003275.

PA (CIBA ) CIBA GEIGY AG.

PI Arvante T;

DR WPI: 1994-214991/26.

PT Aq depot formulations for treatment of e.g. deep vein thrombosis  
 PT - comprises water, hirudin, and a water-soluble salt of calcium,  
 PT magnesium or zinc

XX Disclosure: Page 3-4; 24pp; English.

CC These sequences is a desulphatohirudin variant which lacks the sulphate  
 CC monoester group at Tyr63 of natural hirudin. These proteins have  
 CC approximately the same biological activity as natural, sulphated  
 CC hirudin. These proteins can be used in the depot formulation of the  
 CC invention for the treatment of deep vein thrombosis. The formulations  
 CC comprise water, a hirudin or a hirudin variant and calcium, magnesium  
 CC or zinc ions in the form of water-insoluble salts. These formulations  
 CC have improved stability. When the hirudin is administered using this  
 CC formulation it has been found that there is less bleeding around the  
 CC injection site than when it is administered as a simple solution.

XX Sequence 64 AA:

Query Match 98.6%; Score 363; DB 15; Length 64;  
 Best Local Similarity 100.0%; Pred. No. 5.3e-28;  
 Matches 64; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 LVTYDCTESGONCLCEGSNVCGGKNCILGSDGKNCQVTEGTPKPOSHNDGFEELP 60

DB 1 LVTYDCTESGONCLCEGSNVCGGKNCILGSDGKNCQVTEGTPKPOSHNDGFEELP 60

OY 61 EYLI 64

DB 61 EYLI 64

## RESULT 12

AAP50329  
 ID AAP50329 standard; protein; 65 AA.

AC AAP50329;

DT 12-NOV-1991 (first entry)

DE Hirudin protein.

XX Hirudin; anticoagulant; thrombosis; diagnosis;

XX Hirudo medicinalis.

PN W08504418-A.

PD 10-OCT-1985.

PF 27-MAR-1985; 85WO-FR00062.

PR 27-MAR-1984; 84FR-0004755.

PR 27-APR-1984; 84FR-0013250.

PA (TRAN-) TRANSGENE SA.

PI Tolstoshev P, Harvey R, Courtney M, Lecocq J-P;

DR WPI: 1985-263187/42.

PT Cloning and expression vector contg. DNA for hirudin - or analogues,  
 PT useful as anticoagulant.  
 PS Disclosure: Fig. 1; 92pp; French.

CC DNA encoding hirudin or its analogues can be inserted into cloning  
 CC and expression vectors comprising an origin of replication for  
 CC pBR322, a promoter (esp. all/part of a lambda phage), and an  
 CC initiation region, specifically the sequence ATACACAGACATCTAAG.  
 CC It may also contain all/part of gene N from lambda and/or a gene  
 CC encoding antibiotic resistance. The vector is esp. pTG 720, 718 and  
 CC 719. Hirudin is a known anticoagulant for treating venous  
 CC thrombosis, vascular occlusions or intravenous disseminated  
 CC coagulation. When applied topically it may be used to treat  
 CC haemorrhoids, varicose veins, oedema or psoriasis. Hirudin can also  
 CC be used in extracorporeal blood circulation systems, as a  
 CC diagnostic reagent to detect the form. of clots (when labelled),  
 CC and as an additive to laboratory blood samples. Using the vector  
 CC hirudin can now be produced in large quantities and of consistent  
 CC quality.

XX Sequence 65 AA:

Query Match 97.8%; Score 360; DB 6; Length 65;  
 Best Local Similarity 96.9%; Pred. No. 1e-27;  
 Matches 63; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 1 LVTYDCTESGONCLCEGSNVCGGKNCILGSDGKNCQVTEGTPKPOSHNDGFEELP 60

DB 1 LVTYDCTESGONCLCEGSNVCGGKNCILGSDGKNCQVTEGTPKPOSHNDGFEELP 60

OY 61 EYLIQ 65

DB 61 EYLIQ 65

## RESULT 13

AAP50335  
 ID AAP50335 standard; protein; 65 AA.

AC AAP50335;

DT 12-NOV-1991 (first entry)

DE Hirudin variant.



XX WPT: 1987-164868/24.  
DR N-PSDB; AAN70319.

XX New DNA constructs and hybrid vectors for transformation of yeast  
PT etc. - useful for prodn. and secretion of protein with hirudin  
PT activity for use as thrombin inhibitors.

XX Claim 11; p128; 146pp; English.

XX The preferred DNA construct of the invention contains the PHO5  
CC promoter and a DNA segment consisting of the PHO5 signal sequence  
CC upstream of and in reading frame with a DNA sequence coding for  
CC mature desulphatohirudin. The segment is under the transcriptional  
CC control of the PHO5 promoter and the 3' flanking sequence of the  
CC PHO5 gene.  
CC (Updated on 03-OCT-2002 to add missing OS field.)  
XX

SQ Sequence 65 AA;

Query Match 97.8%; Score 360; DB 8; Length 65;  
Best Local Similarity 96.9%; Pred. No. 1e-27;  
Matches 63; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 1 LVTDTCTESGQNLCTCEGSSNVGGGKNCILGSDGKNCVGTGEGTPKPSHNDGDFEETP 60  
DB 1 VVYTDCTESGQNLCTCEGSSNVGGGKNCILGSDGKNCVGTGEGTPKPSHNDGDFEETP 60

OY 61 EEYLDQ 65  
DB 61 EEYLDQ 65

Search completed: December 30, 2002, 16:16:41  
Job time : 37 secs